

DEADLINE PASSED

Billing Code: 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[PROGRAM ANNOUNCEMENT 00033]

CHILDHOOD LEAD POISONING PREVENTION PROGRAMS (CLPPP)

NOTICE OF AVAILABILITY OF FUNDS

A. Purpose

The Centers for Disease Control and Prevention (CDC) announces the availability of fiscal year (FY) 2000 funds for a cooperative agreement program for new State and competing continuation State and local programs to develop and improve Childhood Lead Poisoning Prevention activities which include building Statewide capacity to conduct surveillance of blood lead levels in children. This program addresses the "Healthy People 2000" priority area of Environmental Health.

The purpose of this program is to provide the impetus for the development, implementation, expansion, and evaluation of State and local childhood lead poisoning prevention program activities which include Statewide surveillance capacity to determine areas at high risk for lead exposure. Also, this cooperative agreement is to carry out the core public health functions of *Assessment, Policy Development, and Assurance* in childhood lead poisoning prevention programs.

Funding for this program will be to:

- 1.) Develop and/or enhance a surveillance system that monitors all blood lead levels.
- 2.) Assure screening of children who are potentially exposed to lead and follow-up care for children who are identified with elevated blood lead levels (BLLs).
- 3.) Assure awareness and action among the general public and affected professionals in relation to preventing childhood lead poisoning.
- 4.) Expand primary prevention of childhood lead poisoning in high-risk areas in collaboration with other government and community-based organizations.

As programs shift emphasis from providing direct screening and follow-up services to the core public health functions, cooperative agreement funds may be used to support and emphasize health department responsibilities to screen high risk children and provide appropriate follow-up services. This includes improving coalitions and partnerships; conducting better and more sophisticated assessments; developing and evaluating policies, program performance, and effectiveness based on established goals and objectives.

B. Eligible Applicants

Applicant eligibility is divided into Part A (New Applicants), Part B (Competing Continuation), and Part C (Supplemental Funding for Alternative Surveillance Assessment/Screening Recommendation Evaluation) defined in the following section.

In the future, CDC plans to shift its program emphasis toward State funding for childhood lead poisoning prevention activities. However, the top five metropolitan statistical areas (SMSAs)/largest cities will be eligible for direct funding for childhood lead poisoning prevention activities indefinitely. **They are New York City, Los Angeles, Chicago, Philadelphia, and Houston.**

1. Part A: Eligible applicants are State health departments or other State health agencies or departments not currently funded by CDC and any eligible SMSA not currently receiving direct funding from CDC for childhood lead poisoning prevention activities. Also eligible are health departments or other official organizational authority (agency or instrumentality) of the District of Columbia, the Commonwealth of Puerto Rico, any territory or possession of the United States, and all federally-recognized Indian tribal governments. **Please note: Local health departments are not eligible to apply for cooperative agreement funding under Part A of this program announcement.**

Applicants encouraged to apply under Part A are: Alaska; Arkansas; Georgia; Hawaii; Idaho; Kansas; Kentucky; Mississippi; Nevada; North Dakota; Oklahoma; South Dakota; Tennessee; Texas and Wyoming.

- 2: Part B: Eligible applicants are those currently funded by the Centers for Disease Control and Prevention whose project

period will expire June 30, 2000. These applicants are:
*Alabama; Arizona; California; Delaware; Detroit ,MI;
Houston, TX; Indiana; Iowa; Maine; Marion County, IN;
Michigan; New Hampshire; Pinellas County, FL; Salt Lake
City, UT; Virginia and Westchester, NY.* In the future, CDC
plans to shift its program emphasis towards State and large
metropolitan statistical areas (SMSAs) which includes
Houston, TX funding for childhood lead poisoning prevention
activities. Consequently, **local applicants eligible for
Part B will only receive funding for a two-year project
period based on satisfactory program performance.** These are
Detroit, MI; Marion County, IN; Pinellas County, FL; Salt
Lake City, UT; and Westchester, NY.

3. **Part C: Eligible applicants are those State applicants
that apply under Part B.** Funding under Part C will only be
considered if the Part B application: (1) is successful and
chosen for funding and (2) has met the program requirement
of submitting data to CDC's national surveillance database.

Additional information for all State applicants

If a State agency applying for grant funds is other than the
official State health department, written concurrence by the
State health department must be provided (for example, the State
environmental health agency).

C. Availability of Funds**Part A: New Applicants**

Up to \$2,500,000 will be available in FY 2000 to fund up to 8 new applicants. CDC anticipates that awards for the first budget year will range from \$75,000 to \$800,000.

Part B: Competing Continuations

Up to \$8,000,000 will be available in FY 2000 to fund up to 17 competing continuation applicants. CDC anticipates that awards for the first budget year will range from \$75,000 to \$1,500,000.

Part C: Supplemental Studies

Up to \$400,000 will be awarded in FY 2000 to fund up to 4 assessment/evaluation studies with a three-year project period. These funds will be awarded to support the development of alternative surveillance assessments and/or to conduct evaluation of the impact of lead screening recommendations. Awards are expected to range from \$70,000 to \$100,000, with the average award being approximately \$85,000. Funds will be awarded for assessment/evaluation studies that address one of the following:

1. Alternative Surveillance Assessment - Assessment of lead exposure in a jurisdictional population or sub-population using an approach to surveillance that differs from the Statewide CBLS system described in this announcement.
2. Screening Recommendation Evaluation - Evaluation of the impact of lead screening recommendations on screening for high-risk children.

Funding for State applicants: To determine the type of program activities and the associated level of funding for an *individual State applicant* for Part A or Part B, please refer to the table below. These are suggested funding guidelines and should not be regarded as absolute funding limits. Addendum 2 in the application package provides an explanation of the factors used to develop categorical funding recommendations. Addendum 3 provides an explanation of the program activities required for each funding category.

Alabama	2	Montana	3
Alaska	3	Nebraska	2
Arizona	3	Nevada	3
Arkansas	2	N. Hampshire	3
California*	1	New Jersey	2
Colorado	3	New Mexico	3
Connecticut	2	New York*	2
Delaware	3	N. Carolina	2
Florida*	3	North Dakota	3
Georgia	2	Ohio	1
Hawaii	3	Oklahoma	2
Idaho	3	Oregon	3
Illinois	1	Pennsylvania	1
Indiana*	3	Rhode Island	2
Iowa	2	S. Carolina	2
Kansas	2	South Dakota	2
Kentucky*	3	Tennessee	2
Louisiana	2	Texas*	1
Maine	3	Utah*	3
Maryland	2	Vermont	3
Mass.	2	Virginia	2
Michigan*	2	Washington	2
Minnesota	2	West Virginia	2
Mississippi	2	Wisconsin	2
Missouri	2	Wyoming	3

* Projected level of effort adjusted to account for currently funded locales.

Funding State Applicants - Part A or Part B: Determine your funding category (Category 1, 2, or 3) according to the table below. The range and average of awards for each funding category follows:

Category 1: \$800,000-\$1,500,000, average award \$1,000,000

Category 2: \$250,000-\$800,000, average award \$520,000

Category 3: \$75,000-\$250,000, average award \$150,000

Awards for Local Applicants (under Part B only): The suggested range of awards for local applicants is \$250,000 to \$450,000.

Additional Information on Funding for all Applicants for Part A, Part B, and Part C New awards are expected to begin on or about July 1, 2000, and are made for 12-month budget periods within project periods not to exceed 2-years for local programs or 3-years for State programs. Estimates outlined above are subject to change based on the actual availability of funds and the scope and quality of applications received. Continuation awards within the project period will be made on the basis of satisfactory progress and availability of funds. *Awards cannot supplant existing funding for CLPP or Supplemental Funding Initiatives.* Funds should be used to enhance the level of expenditures from State, local, and other funding sources.

NOTE:

\$ **Funds may not be expended for medical care and treatment or for environmental remediation of sources of lead exposure.**

However, the applicant must provide a plan to ensure that these program activities are carried out.

\$ Not more than 10 percent (exclusive of Direct Assistance) of any cooperative agreement or contract through the cooperative agreement may be obligated for administrative costs. This 10 percent limitation is in lieu of, and replaces, the indirect cost rate.

D. Program Requirements

SPECIAL REQUIREMENT regarding Medicaid provider status of applicants: Pursuant to section 317A of the Public Health Service Act (42 U.S.C. 247b-1), as amended by Sec. 303 of the "Preventive Health Amendments of 1992" (Public Law 102-531), applicants AND current grantees must meet the following requirements: For CLPP program services which are Medicaid-reimbursable in the applicant's State:

\$ Applicants who directly provide these services must be enrolled with their State Medicaid agency as Medicaid providers.

\$ Providers who enter into agreements with the applicant to provide such services must be enrolled with their State Medicaid agency as providers. An exception to this requirement will be made for providers whose services are provided free of charge and who accept no reimbursement from any third-party payer. Such providers who accept voluntary donations may still be exempted from this requirement.

In order to satisfy this program requirement, please provide a

copy of a Medicaid provider certificate or Statement as proof that you meet this requirement. Failure to include this information would result in your application being returned. Please place this information immediately behind the budget and budget justification pages.

Cooperative Activities

Part A and Part B: New and Competing Continuations

To achieve the purpose of this cooperative agreement program, the recipient will be responsible for the activities listed under **1.**

Recipient Activities and CDC will be responsible for the activities listed under **2. CDC Activities.**

1. Recipient Activities

1. Establish, maintain, or enhance a **Statewide surveillance system** in accordance with CDC guidance. *For local applicants (under Part B),* enhance a data management system that links with the State's surveillance system or develop an automated data management system to collect and maintain laboratory data on the results of blood lead analyses and data on follow-up care for children with elevated BLLs. State recipients should ensure receipt of data from local programs. Local recipients should transfer relevant data to the appropriate State entity in a timely manner for annual submission to CDC.
2. Manage, analyze and interpret individual State surveillance data, and present and disseminate trends

and other important public health findings.

3. Develop, implement and evaluate a Statewide/jurisdiction-wide childhood blood lead screening plan consistent with CDC guidance provided in *Screening Young Children for Lead Poisoning: Guidance for State and Local Public Health Officials*. For local applicants, participate in the Statewide planning process. Make screening recommendations and appropriate local screening strategies available and known to health care providers.
4. Assure appropriate follow-up care is provided for children identified with elevated blood lead levels.
5. Establish effective, well-defined working relationships within public health agencies and with other agencies and organizations at national, State, and community levels (e.g., housing authorities; environmental agencies; maternal and child health programs; State Medicaid Early Periodic Screening, Diagnosis, and Treatment (EPSDT) programs; community and migrant health centers; community-based organizations providing health and social services in or near public housing units, as authorized under Section 330(i) of the PHS Act; State and local epidemiology programs; State and local housing rehabilitation programs; schools of public health and medical schools; and environmental interest groups).
6. For State Programs, provide managerial, technical, analytical, and program evaluation assistance to local

agencies and organizations in developing or strengthening their CLPP programs activities.

2. CDC Activities

1. Provide technical, and scientific assistance and consultation on program development, implementation and operational issues.
2. Provide technical assistance and scientific consultation regarding the development and implementation of all surveillance activities including data collection methods and analysis of data.
3. Assist with data analysis and interpretation of individual State surveillance data and release of national reports. Reports will include analysis of national aggregate data as well as State-specific data.
4. Assist cooperative agreement recipients with communication and coordination among Federal agencies, and other public and private agencies and organizations.
5. Conduct ongoing assessment of program activities to ensure the use of effective and efficient implementation strategies.

Part C: Supplemental Studies

To achieve the purpose of this program, the recipient will be responsible for the activities listed under **1. Recipient Activities** and CDC will be responsible for the activities listed under **2. CDC Activities**.

1. Recipient Activities

1. Develop and implement a study protocol to include the following: methodology, sample selection, field operation, and statistical analysis. Applicants must provide a means of assuring that the results of the study will be published.
2. Revise, refine, and carry out the proposed methodology for conducting *Supplemental Funding Studies*.
3. Monitor and evaluate all aspects of the assessment activities.
4. Conduct and evaluate public health programs and/or have access to professionals who are knowledgeable in conducting such activities.

2. CDC Activities

1. Provide technical and scientific consultation on activities related to overall program requirements of supplemental funding activities.
2. Provide technical assistance to program manager and/or principal investigator regarding revision, refinement, and implementation of study design and proposed methodology for conducting supplemental funding activities.
3. Assist program manager and/or principal investigator with data interpretation and analysis issues.

E. Application Content

Use the information in the *Program Requirements, Other Requirements, and Evaluation Criteria* sections to develop the application content. Your application will be evaluated on the criteria listed, so it is important to follow them in laying out your program plan:

- # Applications must be developed in accordance with PHS Form 5161-1.
- # Part B applicants also competing for Part C funds must submit separate applications.
- # Application pages must be clearly numbered, and a complete index to the application and its appendices must be included.
- # The original and two copies of the application set must be submitted UNSTAPLED and UNBOUND. All material must be typewritten, double spaced, printed on one side only, with un-reduced font (10 or 12 point font only) on 8-1/2" by 11" paper, and at least 1" margins and heading and footers. All graphics, maps, overlays, etc., should be in black and white and meet the above criteria.
- # **A one-page, single-spaced, typed abstract must be submitted with the application. The heading should include the title of the program, project title, organization, name and address, project director, telephone number, facsimile number, and e-mail address.**
- # **The main body of the CLPP program application (Parts A or B) must include the following: budget/budget justification; Medicaid certification; progress report (Part B applicants only); understanding the problem; surveillance/data-**

management activities; Statewide/jurisdiction-wide planning and collaboration; core public health functions; goals and objectives; program management and staffing; and program evaluation.

- # The main body of the supplemental funding project application (Part C) must include the following: study protocol, project personnel, and project management.
- # Each application should not exceed 75 pages. The abstract, budget narrative, and budget justification pages are not included in the 75 page limit. Supplemental information should be placed in appendices and is not to exceed 25 pages.
- # Part B applicants must submit a progress report no longer than 10 pages in their competing continuation application. **This report should be placed immediately after the budget and budget justification.**
- # Provide qualified staff, other resources, and knowledge to implement the provisions of the program. Applicants requesting cooperative agreement supported positions must provide assurances that such positions will be authorized to be filled by the applicant-s personnel system.

F. Submission and Deadline

Submit the original and two copies of the PHS 5161-1 (OMB Number 0937-0189) on or before April 12, 2000. Forms are in the application kit.

Submit the application to:

Mattie B. Jackson, Grants Management Specialist

Grants Management Branch, Procurement and Grants Office

Program Announcement 00033

Centers for Disease Control and Prevention (CDC)

2920 Brandywine Road, Room 3000

Atlanta, GA 30341-4146

Applications shall be considered as meeting the deadline if they are either: (1) received on or before the deadline date, or (2) sent on or before the deadline date and received in time for submission for the review process. Applicants must request a legibly dated receipt from a commercial carrier or U. S Postal Service. Private metered postmarks shall not be acceptable as proof of timely mailing.

Applications which do not meet the criteria above are considered late applications. Late applications will not be considered in the current competition and will be returned to the applicant.

G. Evaluation Criteria

The review of applications will be conducted by an objective review panel as they relate to the applicant's response to either Part A, Part B, or Part C. The applications will be evaluated according to the following criteria:

PART A: New Applicants

1. Understanding of the Problem (15 points)

The extent to which the applicant's description and understanding of the burden and distribution of childhood lead exposure or elevated BLLs in their jurisdiction, using evidence (as available) of incidence and/or prevalence and demographic indicators, including a description of the Medicaid population.

2. Surveillance Activities (20 points)

The applicant's ability to develop a childhood blood lead surveillance system that includes; (a) a flow chart that describes data transfer, (b) a mechanism for tracking lead screening services to children, especially Medicaid children, and (c) a mechanism for reporting data annually to the CDC's national surveillance database. The clarity, feasibility, and scientific soundness of the surveillance approach. Also, the extent to which the proposed time table for accomplishing each activity and methods for evaluating each activity are appropriate and clearly defined. The following elements will be specifically evaluated:

1. How laboratories report Blood Lead Levels (BLLs), including ability to identify and assure reporting from private laboratories which perform lead testing.
- b. How data will be collected and managed.
3. How quality of data and completeness of reporting will be assured.
4. How and when data will be analyzed.
- e. How summary data will be reported and disseminated on a regular basis (i.e., newsletters, fact sheets, annual reports).

6. Protocols for follow-up of individuals with elevated BLLs.
 - g. Provisions to obtain denominator data (results of all laboratory blood lead tests, regardless of level).
 8. Time line and methods for evaluating the Childhood Blood Lead Surveillance (CBLs) approach.
 9. Plans to convert paper-based components of the system to electronic data manipulation.
 10. Use of data including evaluation of prevention activities, especially to target screening and prevention efforts.
3. Statewide Planning and Collaboration (20 points)
- The applicant-s ability to develop Statewide screening recommendations with appropriate local strategies. The following elements will be specifically evaluated:
1. The proposed approach to developing and carrying out an inclusive State- wide screening plan as outlined in *Screening Young Children for Lead Poisoning: Guidance for State and Local Health Officials*.
 2. The extent to which the applicant plans to utilize surveillance and program data to produce a Statewide screening recommendation, with specific attention given to the Medicaid population.
 3. The ability of the applicant to involve collaborators in the development of a screening plan and implementation of strategies to strengthen childhood lead poisoning prevention activities.
 4. The applicant-s demonstrated ability to collaborate

with principal partners, including managed-care organizations, State Medicaid agency, child health-care providers and provider groups, insurers, community-based organizations, housing agencies, and banking, real-eState, and property-owner interests, must be demonstrated by letters of support, memoranda of understanding, contracts, or other documented evidence of relationships.

4. Capacity to Carry out Public Health Core Functions (10 points)

The applicant's ability to describe the approach and activities necessary to achieve a balance among health department roles in CLPP, including assessment, program and policy development, and monitoring, evaluating, and ensuring the provision of all necessary components of a comprehensive CLPP activities within their respective categories.

5. Goals and objectives (15 points)

The extent to which the applicant's goals and objectives relate to the CLPP activities in their respective categories. Objectives must be relevant, specific, measurable, achievable, and time-framed. There must be a formal work plan with a description of methods, a timetable and program staff responsible for accomplishment of each objective, and the evaluation of each proposed objective.

6. Project management and staffing (10 points)

The extent to which the applicant has documented the skills

and ability to develop and carry out a comprehensive CLLP program. Specifically, the applicant should:

1. Describe the proposed health department staff roles in CLPP, their specific responsibilities, and their level of effort and time. Include a plan to expedite filling of all positions and assure that requested positions have been or will be approved by applicant's personnel system.
2. Describe the plan to provide training and technical assistance to health department personnel and consultation to collaborators outside the health department, including proposed design of information-sharing systems.

7. Program evaluation (10 points)

The extent to which the applicant proposes to measure the overall impact of health department CLPP activities.

Specific criteria should include:

- a. The plan for evaluating the impact or outcome of CLPP activities, including evaluation design, methods, and activities.
2. Description of how the project will assess changes in public policy and measure the effectiveness of collaborative activities.
3. Progress made in childhood lead poisoning prevention which resulted from planned health department strategies.

8. Budget justification (not scored)

The extent to which the budget is reasonable, clearly

justified, and consistent with the intended use of funds.

PART B: Competing Continuations

1. Understanding of the Problem (15 points)

The extent to which the applicant's description and understanding of the burden and distribution of childhood lead exposure or elevated BLLs in their jurisdiction, using evidence (as available) of incidence and/or prevalence and demographic indicators, including a description of the Medicaid population.

2. Surveillance activity (20 points)

For State Applicants: The applicant's ability to expand its childhood blood lead surveillance system that includes tracking lead screening for Medicaid children, evaluating the existing system, and reporting data to the CDC's national surveillance database. The clarity, feasibility, and scientific soundness of the surveillance approach. Also, the extent to which the proposed time table for accomplishing each activity are appropriate and clearly defined. The following elements will be specifically evaluated:

- a. How laboratories report BLLs, including ability to identify and assure reporting from private laboratories which perform lead testing.
- b. How data are collected and managed.
3. How quality of data and completeness of reporting are assured.
4. How and when data are analyzed.

- e. How summary data are reported and disseminated on a regular basis (i.e., newsletters, fact sheets, annual reports).
6. Protocols for follow-up of individuals with elevated BLLs.
- g. Provisions to obtain denominator data (results of all laboratory blood lead tests, regardless of level).
8. Time line and methods for evaluating the Childhood Blood Lead Surveillance (CBLs) approach.
9. Process used to convert paper-based components of the system to electronic data.
10. Use of data including evaluation of prevention activities, especially to target screening and prevention efforts.

For local applicants (Part B only): The applicant's ability to expand their data management system, including the approach to participating in the State CBLs, if applicable. The clarity, feasibility, and scientific soundness of the approach to data management. Also, the extent to which the proposed schedule for accomplishing each activity and method for evaluating each activity are clearly defined and appropriate. The following elements will be specifically evaluated:

- a. How laboratories report Blood Lead Levels (BLL), including ability to identify and assure reporting from private laboratories which perform lead testing.
- b. How data are collected and managed.
- c. How quality of data and completeness of reporting are assured.

- d. How and when data are analyzed.
 - e. How summary data are reported and disseminated on a regular basis (i.e., newsletters, fact sheets, annual reports).
 6. Protocols for follow-up of individuals with elevated BLLs.
 - g. Provisions to obtain denominator data (results of all laboratory blood lead tests, regardless of level).
 - h. Time line and methods for evaluating the Childhood Blood Lead Surveillance (CBLs) approach.
 - i. Process used to convert paper-based components of the system to electronic data.
 - j. Use of data including evaluation of prevention activities, especially to target screening and prevention efforts.
3. Statewide/Jurisdiction-wide Planning and Collaboration (20 points)

The applicant's ability to develop Statewide/jurisdiction-wide screening recommendations with appropriate local strategies. The following elements will be specifically evaluated:

1. The approach to developing and carrying out an inclusive State- or jurisdiction-wide screening plan as outlined in *Screening Young Children for Lead Poisoning: Guidance for State and Local Health Officials*.
2. The extent to which the applicant utilized surveillance and program data to produce Statewide/jurisdiction-wide

screening recommendations and target the Medicaid population.

3. Description of how collaborations facilitated the development of a screening plan and strengthened childhood lead poisoning prevention strategies.
4. Evidence of collaboration with principal partners, including managed-care organizations, State Medicaid agency, child health-care providers and provider groups, insurers, community-based organizations, housing agencies, and banking, real-eState, and property-owner interests. These collaborations must be demonstrated by letters of support, memoranda of understanding, contracts, or other documented evidence of relationships.

Note: For applicants under Part B, describe progress in developing and implementing the screening plan based upon each of the elements listed above.

4. Capacity to carry out public-health core functions (10 points)

The ability to describe the approach and activities taken to achieve a balance among health-department roles in CLPP, including assessment, program and policy development, and monitoring, evaluating, and ensuring the provision of all CLPP activities within their respective categories (for example, Category 3 requires screening plans, surveillance systems, assure follow-up care, and evaluation).

5. Goals and objectives (10 points)

The extent to which the applicant's goals and objectives relate to the CLPP activities in their respective categories under which they applied. Objectives must be relevant, specific, measurable, achievable, and time-framed. There must be a formal work plan with a description of methods and a timetable and program staff responsible for accomplishment of each objective.

6. Project management and staffing (10 points)

The extent to which the applicant has the skills and ability to develop and carry out a comprehensive CLLP program.

Specifically the applicant should:

1. Describe the proposed health department staff roles in CLPP, their specific responsibilities, and their level of effort and time. Include a plan to expedite filling of all positions and assure that requested positions have been or will be approved by the applicant's personnel system.
2. Describe the plan to provide training and technical assistance to health department personnel and consultation to collaborators outside the health department, including proposed design of information-sharing systems.

7. Program evaluation (10 points)

The extent to which the applicant measures the overall impact of health department CLPP activities. Specific criteria should include:

- a. Description of the progress made to evaluate the impact

and outcome of collective CLPP activities, including the evaluation design, methods, and tasks.

2. Description of the changes in the effectiveness of collaborative activities.
3. Progress made in childhood lead poisoning prevention which resulted from planned health department strategies.

8. Budget justification (not scored)

The extent to which the budget is reasonable, clearly justified, and consistent with the intended use of funds.

PART C: SUPPLEMENTAL FUNDING - Factors to be Considered

1. Study protocol (45 points)

The applicant's ability to develop a scientifically sound

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2. Project personnel (20 points)

The extent to which personnel involved in this project are qualified, including experience in conducting relevant studies. In addition, the applicant's ability to commit appropriate staff time needed to carry out the study.

3. **Project management (35 points)**

The applicant's ability to implement and monitor the proposed study to include specific, attainable, and realistic goals and objectives, and evaluation plan.

4. Budget justification (not scored)

The extent to which the budget is reasonable, clearly justified, and consistent with the intended use of cooperative agreement funds.

5. **Human subjects (not scored)**

The extent to which the applicant complies with the Department of Health and Human Services regulations (45 CFR Part 46) on the protection of human subjects.

H. Other Requirements

Technical Reporting Requirements

Provide CDC with the original plus two copies of:

1. Quarterly progress reports, which are required of all grantees. **The quarterly report narrative should not exceed 25 pages.** Time lines for the quarterly reports will be established at the time of award, but are typically due 30 days after the end of each quarter.
2. Calendar year surveillance data and a written surveillance

summary report must be submitted annually to CDC in the approved OMB format to be disseminated to State and local public health officials and congressional personnel. Data must be submitted to CDC by March 31st in the required format for analysis.

3. Financial Status Reports, are due within 90 days of the end of the budget period.
4. Final financial reports and performance reports are due within 90 days after the end of the project period.

Send all reports to the Grants Management Specialist identified in the [Where to Obtain Additional Information](#) section of this announcement.

NOTE: Data collection initiated under this cooperative agreement program has been approved by the Office of Management and Budget under OMB number (0920-0337), [National Childhood Blood Lead Surveillance System](#), Expiration Date: March 31, 2001.

The following additional requirements are applicable to this program. For a complete description of each, see Addendum 1 in the application package.

- AR-1 Human Subjects Requirement
- AR-2 Requirements for Inclusion of Women and Racial and Ethnic Minorities in Research
- AR-7 Executive Order 12372 Review
- AR-9 Paperwork Reduction Act Requirements

- AR-10 Smoke-Free Workplace Requirements
- AR-11 Healthy People 2000
- AR-12 Lobbying Restrictions

I. Authority and Catalog of Federal Domestic Assistance Number

This program is authorized under sections 301(a), 317A and 317B of the Public Health Service Act [42 U.S.C. 241(a), 247b-1, and 247b-3], as amended. Program regulations are set forth in Title 42, Code of Federal Regulations, Part 51b. The Catalog of Federal Domestic Assistance number is 93.197.

J. Pre-Application Workshop for New and Competing Continuation Applicants

In addition, for interested applicants, a telephone conference call for pre-application technical assistance will be held on Wednesday, February 16, 2000, from 1:30 p.m. to 3:30 p.m, Eastern Standard Time. **The bridge number for the conference call is 1-800-311-3437, and the pass code is 350892.** For further information about all workshops, please contact Claudette Grant-Joseph at 404-639-2510.

11. Where to Obtain Additional Information:

This and other CDC announcements may be downloaded through the CDC homepage on the Internet at <http://www.cdc.gov>. Please refer to program announcement number 00033 when requesting information.

To receive additional written information and to request an application kit, call 1-888-GRANTS4 (1-888-472-6874). You will be asked to leave your name, address, and phone number and will

need to refer to Announcement 00033. You will receive a complete program description, information on application procedures, and application forms. CDC will not send application kits by facsimile or express mail.

If you have questions after reviewing the contents of all documents, business management technical assistance may be obtained from:

Mattie B. Jackson, Grants Management Specialist
Grants Management Branch, Procurement and Grants Office
Centers for Disease Control and Prevention (CDC)
2920 Brandywine Road, Room 3000
Atlanta, GA 30341-4146
telephone (770) 488-2718
Internet address **mij3@cdc.gov**

For programmatic technical assistance, contact:

Claudette A. Grant-Joseph, Chief,
Program Services Section, Lead Poisoning Prevention Branch
Division of Environmental Hazards and Health Effects
National Center for Environmental Health
Centers for Disease Control and Prevention (CDC)
1600 Clifton Road, NE, Mailstop E-25
Atlanta, GA 30333
telephone (404) 639-2510
Internet address **cag4@cdc.gov**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
Program Announcement 00033
Childhood Lead Poisoning Prevention Programs**

AR-1

Human Subjects Requirements

If the proposed project involves research on human participants, the applicant must comply with the Department of Health and Human Services Regulations (45 CFR 46) regarding the protection of human research participants. Assurance must be provided to demonstrate that the project will be subject to initial and continuing reviews by an appropriate institutional review board. The applicant will be responsible for providing evidence of this assurance in accordance with the appropriate guidelines and forms provided in the application kit.

In addition to other applicable committees, Indian Health Service (IHS) institutional review committees also must review the project if any component of IHS will be involved with or will support the research. If any American Indian community is involved, its tribal government must also approve that portion of the project applicable to it.

Unless the awardee holds a Multiple Project Assurance, a Single Project Assurance is required, as well as an assurance for each subcontractor or cooperating institution that has immediate responsibility for human participants.

The Office for Protection from Research Risks (OPRR) at the National Institutes of Health (NIH) negotiates assurances for all activities involving human participants that are supported by the Department of Health and Human Services.

AR-2

Requirements for Inclusion of Women and Racial and Ethnic Minorities in Research

It is the policy of the Centers for Disease Control and Prevention (CDC) and the Agency for Toxic Substances and Disease Registry (ATSDR) to ensure that individuals of both sexes and the various racial and ethnic groups will be included in CDC/ATSDR-supported research projects involving human subjects, whenever feasible and appropriate. Racial and ethnic groups are those defined in OMB Directive No. 15 and include American Indian or Alaska Native, Asian, Black or African American, Hispanic or Latino, Native Hawaiian or Other Pacific Islander. Applicants shall ensure that women, racial and ethnic minority populations are appropriately represented in applications for research

involving human subjects. Where clear and compelling rationale exist that inclusion is inappropriate or not feasible, this situation must be explained as part of the application. This policy does not apply to research studies when the investigator cannot control the race, ethnicity, and/or sex of subjects. Further guidance to this policy is contained in the Federal Register, Vol. 60, No. 179, pages 47947-47951, and dated Friday, September 15, 1995.

AR-7

Executive Order 12372 Review

Applications are subject to Intergovernmental Review of Federal Programs, as governed by Executive Order (E.O.) 12372. The order sets up a system for State and local governmental review of proposed Federal assistance applications. Applicants should contact their State single point of contact (SPOC) as early as possible to alert the SPOC to prospective applications and to receive instructions on the State process. For proposed projects serving more than one State, the applicant is advised to contact the SPOC for each State affected. (The application kit contains a current list of SPOCs.) SPOCs who have recommendations about the State process for applications submitted to CDC should send them, in a document bearing the program announcement number, no more than 60 days after the application deadline date, to:

Mattie B. Jackson, Grants Management Specialist
Grants Management Branch, Procurement and Grants Office
Announcement Number 00033
Centers for Disease Control and Prevention
2920 Brandywine Road, Room 3000
Atlanta, GA 30341

Indian tribes must request tribal government review of their applications.

If Indian tribes are eligible for the program, change the sentence about SPOC recommendations as follows:

SPOCs or tribal governments that have recommendations about an application submitted to CDC should send them, in a document bearing the program announcement number, no more than 60 days after the application deadline date, to:

Mattie B. Jackson, Grants Management Specialist
Grants Management Branch, Procurement and Grants Office
Announcement Number 00033
Centers for Disease Control and Prevention
2920 Brandywine Road, Room 3000
Atlanta, GA 30341

CDC does not guarantee to accept or justify its nonacceptance of recommendations that are received more than 60 days after the

application deadline.

AR-9

Paperwork Reduction Act Requirements

Projects that involve data collection from 10 or more persons and that are funded by grants and cooperative agreements will be subject to review and approval by the Office of Management and Budget (OMB).

Data collection initiated under this grant/cooperative agreement) has been approved by the Office of Management and Budget (OMB) under OMB number 0920-0337 for CDC), National Childhood Blood Lead Surveillance System, expiration date March 31, 2001.

AR-10

Smoke-Free Workplace Requirements

CDC strongly encourages all recipients to provide a smoke-free workplace and to promote abstinence from all tobacco products. Public Law 103-227, the Pro-Children Act of 1994, prohibits smoking in certain facilities that receive Federal funds in which education, library, day care, health care, or early childhood development services are provided to children.

AR-11

Healthy People 2000

CDC is committed to achieving the health promotion and disease prevention objectives of A Healthy People 2000, @ a national activity to reduce morbidity and mortality and improve the quality of life. For a copy of "Healthy People 2000" (Full Report:

Stock No. 017-001-00474-0) or "Healthy People 2000" (Summary Report: Stock No. 017-001-00473-1), write or call:

Superintendent of Documents

Government Printing Office

Washington, DC 20402-9325

Telephone (202) 512-1800

AR-12

Lobbying Restrictions

Applicants should be aware of restrictions on the use of HHS funds for lobbying of Federal or State legislative bodies. Under the provisions of 31 U.S.C. Section 1352, recipients (and their sub-tier contractors) are prohibited from using appropriated Federal funds (other than profits from a Federal contract) for lobbying congress or any Federal agency in connection with the award of a particular contract, grant, cooperative agreement, or loan. This includes grants/cooperative agreements that, in whole or in part, involve conferences for which Federal funds cannot be used directly or indirectly to encourage participants to lobby or to instruct participants on how to lobby.

In addition no part of CDC appropriated funds, shall be used, other than for normal and recognized executive-legislative relationships, for publicity or propaganda purposes, for the preparation, distribution, or use of any kit, pamphlet, booklet, publication, radio, television, or video presentation designed to support or defeat legislation pending before the Congress or any State or local legislature, except in presentation to the Congress or any State or local legislature itself. No part of the appropriated funds shall be used to pay the salary or expenses of any grant or contract recipient, or agent acting for such recipient, related to any activity designed to influence legislation or appropriations pending before the Congress or any State or local legislature.

Addendum 2

Background on CDC's estimate of number and proportion of children at high risk for lead exposure by State

To provide States with general guidance about the appropriate amount of funding to request under this Program Announcement, CDC estimated the number and percentage of children with elevated BLLs for each State. CDC used a logistic-regression model to estimate the contribution of four major risk factors to the probability that an individual child would have a blood lead level (BLL) of at least 10 µg/dL. The selected risk factors were based on data from Phase 2 of the Third National Health and Nutrition Examination Survey (NHANES III, Phase 2) and included the age and race of children, age of housing, and family income.

The model established a relative contribution or Acoefficient® for each of these factors. These coefficients were then applied to the relevant categories of 1990 census data for each State to produce an estimate of both *the number* and *the percentage* of children with elevated BLLs in the State.

CDC's purpose in estimating the number and percentage of children with EBLs in each State is to approximate the level of effort that may be required to provide prevention services to the entire population of a State. In accordance with this purpose, CDC adjusted the level of effort projected for State-level CLPP Programs in States with one or more locales currently receiving separate funding under this grant program.

To derive the funding category for each State, CDC gave twice as much weight to the estimated percentage of children with elevated BLLs as to the estimated number of children with elevated BLLs.

Note 1: The categorization scheme developed for use in this Program Announcement is likely to be of only limited usefulness for other purposes. The use of an approximation is necessary because of the wide variation among States in the extent to which their pediatric populations are exposed to lead.

Note 2: Applicants are encouraged to use the funding category that is suggested for the applicant's State; however, note these are suggested funding guidelines and should not be regarded as absolute funding limits.

Addendum 3

Description of Program Activities by Funding Categories

Funding levels are associated with category type and level of program activity to be supported by CDC. However, **regardless of category type**, all programs are required to develop and implement screening plans and have a surveillance system designed to monitor blood lead levels in children. Following are the minimum requirements for each category.

At a minimum:

Category 1 applicants are to use CDC funding to: implement and evaluate screening plans; submit and analyze data from a Statewide surveillance system; assure screening and follow-up care; provide public and professional health education and health communication; conduct program impact evaluation; and implement primary prevention activities.

Category 2 applicants are to use CDC funding to: implement and evaluate screening plans; submit and analyze data from a Statewide surveillance system; assure screening and follow-up care; provide public and professional health education and health communication; and conduct program impact evaluation.

Category 3 applicants are to use CDC funding to: implement and evaluate screening plans; submit and analyze data from a Statewide surveillance system; assure screening and follow-up care; and conduct program impact evaluation.

Addendum 4

BACKGROUND AND DEFINITIONS

Background:

In the last few years, there have been three major changes in the context within which CLPP and CBLIS programs function. These are:

- \$ **Changing functions of health departments.** Many health departments have ceased to be major providers of direct screening and follow-up care services, as Medicaid beneficiaries who formerly received preventive health care in health departments have enrolled in managed-care organizations. A decrease in funding has occurred in many health departments.
- \$ **Renewed emphasis on accountability of government agencies.** A renewed call for accountability in government agencies requires that health departments document both the need for and the impact of their programs.
- \$ **Continuing declines in BLLs of the entire U.S. population, resulting in wide variation among jurisdictions with regard to the magnitude of their childhood lead poisoning problems.**

Resource limitations and the demand for public accountability have made it increasingly important for health departments to perform the core functions of public health as outlined in *The Future of Public Health* (IOM, 1988). These core functions are assessment, policy development, and assurance. Health department personnel must also accomplish their missions through others, by deepening relationships among new and old partners both in and outside of the health department. Also, the widening disparity among jurisdictions with regard to the magnitude of the childhood lead poisoning problem has focused attention on State and local health departments, as opposed to the Federal government, as the appropriate decision-makers for lead screening. Taken together, these changes are having a profound impact on CLPP programs, necessitating a change in programmatic emphasis.

CLPP and CBLIS programs are positioned to bring about improved screening and follow-up care for children with elevated BLLs, improved public and professional awareness of the problem of childhood lead poisoning, and improved childhood blood lead surveillance, by performing the three core public health functions related to childhood lead poisoning prevention.

Definitions

- \$ *Assessment*: Activities organized by a health department for the purpose of determining the risk for lead exposure among the children in its jurisdiction and the adequacy of programmatic activities to address this risk.
- \$ *Assurance*: Activities organized by a health department for the purpose of 1) monitoring the provision of CLPP services including screening, follow-up care, and public and professional education; and 2) ensuring, as a provider of last resort, the availability of necessary services.
- \$ *Care coordination*: The monitoring and organizing of follow-up care for a child with an elevated blood lead level (BLL). Follow-up care includes both medical and environmental interventions.
- \$ *High-risk*: A term used to designate areas, populations, and individuals with risk for lead exposure that is assessed or demonstrated to be higher than average.
- \$ *Lead hazard*: Accessible paint, dust, soil, water, or other source or pathway that contains lead or lead compounds that can contribute to or cause elevated BLLs.
- \$ *Lead hazard remediation*: The elimination, reduction, or containment of known and accessible lead sources.
- \$ *Policy development*: Activities organized by a health department for the purpose of framing the CLPP problem and establishing the response to it in its jurisdictions; includes development, oversight, and evaluation of necessary programs, relationships, and policies that will support CLPP.
- \$ *Primary prevention*: The prevention of elevated BLLs in an individual or population, usually by reducing or eliminating lead hazards in the environment.
- \$ *Program*: A designated unit within an agency responsible for implementing and coordinating a systematic and comprehensive approach to CLPP and CBLS.
- \$ *Surveillance*: A process which 1) systematically collects information over time about children with elevated BLLs using laboratory reports as the data source; 2) provides for the follow-up of cases, including field investigations when necessary; 3) provides timely and useful analysis and reporting of the accumulated data, including an estimate of the rate of elevated BLLs among all children receiving blood tests; and 4) reports data to CDC in the appropriate format.

Addendum 5

Childhood Lead Poisoning Prevention Program Components

- Component 1. Statewide/Jurisdiction-wide Screening Plan (Required activity for all funded applicants).** Development of a Statewide CBLS (childhood blood lead screening) plan consistent with CDC guidance provided in *Screening Young Children for Lead Poisoning: Guidance for State and Local Public Health Officials*.
- Component 2. Statewide Surveillance System (Required activity for all funded State applicants).** Development of a CBLS system that includes collection, analysis, and dissemination of data on: screening, prevalence of elevated BLLs, sources of lead exposure, and follow-up care among children. Inclusion of surveillance data in the national CBLS database maintained by CDC. [Funded locales also need to engage in planning, data management, and surveillance, but it is likely that these activities will take place within the context of State activities.]
- Component 3. Assurance of screening and follow-up care (Required activity for all funded applicants).** Development, improvement, and oversight of lead-related policies and services associated with: a) screening; b) follow-up care for those with elevated BLLs, including care coordination, family education about lead exposure, and environmental investigation; and c) remediation of lead hazards. *Of particular interest are efforts to develop policies and to convene and coordinate concerned and responsible parties to bring about these activities.*
- Component 4. Public and professional health education and health communication (Required for Funding Categories 1 & 2)** Development, improvement, and oversight of strategies to perform health education and health communication about CLPP for a variety of target audiences. [Note: The ability to communicate CLPP program goals effectively and to educate community members about CLPP underlie all other aspects of the CLPP program.]

Component 5. Evaluation of program impact (Required activity for all funded applicants).
Monitoring and evaluation of the effectiveness of screening, follow-up, education and communication, lead-hazard remediation, and primary prevention activities to ensure that programs are consistent with plans and policies, and revision of programmatic efforts as necessary on the basis of evaluation findings. (For example: What is your program's expected outcome as a result of all program activities implemented)

Component 6. Primary prevention (Required for Funding Category 1). Development, improvement, and oversight of policies and strategies to bring about primary prevention.

Provide a work plan for health department activities in each of their respective CLPP program components listed above. The work plan should outline long-range goals for the duration of the project period. You must include detailed, specific, measurable, and time-phased objectives for the first budget year of the proposed project. Include a tentative work plan and time line for the remaining years of the proposed project.

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